LATVIJAS VIDES, ĢEOLOĢIJAS UN METEOROLOĢIJAS CENTRS

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BERKEM DEVELOPPEMENT S.A.

Le Marais Ouest
FR 24680, Gardonne
France

On authorisation of the biocidal product AXIL 3000 P+ through mutual recognition in Latvia

Latvian Environment, Geology and Meteorology Centre (LEGMC) has evaluated an application submitted by BERKEM DEVELOPPEMENT S.A. on 8th September 2020 concerning an authorisation of the biocidal product *AXIL 3000 P+* through mutual recognition in Latvia.

LEGMC has agreed with Product Assessment Report and Summary of Product Characteristics for *AXIL 3000 P+* developed by the reference Member State – Belgium.

Therefore, in accordance with Article 33 of *Regulation (EU) No 528/2012 of the European Parliament and of the Council of 22 May 2012 concerning the making available on the market and use of biocidal products* (regulation 528/2012), LEGMC authorises the biocidal product *AXIL 3000 P+*.

AXIL 3000 P+ contains:

- 1.0% of *3-iodo-2-propynyl butylcarbamate* (CAS No.55406-53-6, EC No.259-627-5),
- 1.1% of *propiconazole* (CAS No. 60207-90-1, EC No. 262-104-4),
- 1.1% of *tebuconazole* (CAS No. 107534-96-3, EC No. 403-640-2) and
- 2.0% of *permethrine* (CAS No. 52645-53-12, EC No. 258-067-9).

LEGMC assigns the authorisation number **LV/2020/MR/017** for **AXIL 3000 P+**.

The authorisation number is valid until **18 April 2024.**

The authorisation number shall be indicated on the label of the biocidal product.

The authorisation of the *AXIL 3000 P+* is granted on the following terms:

- Product type: 8 – Wood preservative;
- Target organisms: wood-boring insects, termites and wood destroying fungi (white and brown rot fungi);
- Fields of use: Use Class 1, 2 and 3 (fully automated dipping, fully automated spraying, vacuum pressure (autoclave));

- Users: professionals;
- Product description: micro emulsion;
- Pack sizes and packaging material: Can/Tin, HDPE (25L); Drum, HDPE (60L); Drum, HDPE (220L); IBC, HDPE (640L); IBC, HDPE (1000L);
- Product stability: up to 24 months.

The authorisation applies only to the *AXIL 3000 P+* in the composition, form and packing for which the first authorisation is granted by reference Member State.

The information on the label (and if applicable an enclosed instruction of use) of the *AXIL 3000 P+* shall be as it is indicated in the first authorisation of above mentioned product, taking into account also the information which is stated in the Product Assessment Report and Summary of Product Characteristics issued by reference Member State.

The information on the label shall be in Latvian.

Notwithstanding content of the label specified above, requirements stated in:

- Article 69 Regulation (EU) No 528/2012 of the European Parliament and of the Council of 22 May 2012 concerning the making available on the market and use of biocidal products;
- Regulation (EC) No. 1272/2008 of the European Parliament and of the Council of 16 December 2008 on classification, labelling and packaging of the substances and mixtures, amending and repealing Directives 67/548/EEC and 1999/45/EC, and amending Regulation (EC) No 1907/2006;
- all other relevant legislation shall be applied.

BERKEM DEVELOPPEMENT S.A. shall inform LEGMC about any changes in accordance with *Commission Implementing Regulation (EU) No 354/2013 of 18th April 2013 on changes of biocidal products authorised in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council.*

If the first authorisation issued by reference Member State is amended or revoked, the authorisation of *AXIL 3000 P+* may be re-opened for review before 18 April 2024.

Additionally, LEGMC would like to inform that BERKEM DEVELOPPEMENT S.A. is fully responsible of the content of the biocidal product *AXIL 3000 P+,* as well as its classification, labelling, instruction of use and safety data sheet.

LEGMC would like to ask BERKEM DEVELOPPEMENT S.A. to notify the above mentioned information down to supply chain.

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